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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

US Pat No. 6,846,807; Issued Jan 25, 2005 Customer No. 23379

Serial No. 10/627,451; Filed: July 25, 2003 Confirmation No. 3438

Inventor: Graff et al Group Art Unit:1623

Docket No. UTSD:0980 Examiner:Khare, Devesh

Title: *Colorectal Neoplasia*

CERTIFICATE OF TRANSMISSION

I hereby certify that this corr is being transmitted by facsimile to the Commr for Patents 703-872-9306 on March 3, 2005.

Signed


Richard Aron OsmanREQUEST FOR CERTIFICATE OF CORRECTION UNDER 37 CFR 1.322

Mail Stop CERTIFICATE OF CORRECTION
Commissioner for Patents
P.O. Box 1450
Alexandria, VA 22313-1450

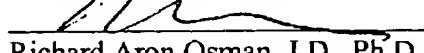
Dear Commissioner:

The Assignee of this Patent requests that the Commissioner issue a certificate of correction in this Patent. The Office erroneously neglected to print the authorized Examiner's amendments in the Notice of Allowability dated Oct 21, 2004 (p.2-3, attached). In particular, the Examiner amended claims 5, 6, 12 and 13 (renumbered from original claims 5, 7, 13 and 14) as shown on the attached PTO/SB44.

Accordingly, please correct the Patent cover sheet as indicated on the attached Form PTO/SB/44. This correction includes no new matter.

Although an Office oversight introduced the error in the issued Patent, as a procedural precaution, Applicants alternatively submit this request under 37CFR1.323. The Commissioner is authorized to charge any fees or credit any overpayments associated with this communication to our Deposit Account No. 19-0750 (order no. UTSD:0980).

Respectfully submitted,
SCIENCE & TECHNOLOGY LAW GROUP


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enc. Notice Allowability dated 10-21-04, p.2-3 (2p).
PTO/SB44 (1p)

PTO/SB/44 (04-04)

Approved for use through 04/30/2007. OMB 3651-0033
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(Also Form PTO-1050)

UNITED STATES PATENT AND TRADEMARK OFFICE
CERTIFICATE OF CORRECTION

PATENT NO. : 6,846,807

DATED : Jan 25, 2005

INVENTOR(S) : Jonathan Graff & Matthew Wieduwilt

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

In claim 1, line 4, the "determining" step should read:
determining a patient is subject to colorectal neoplasia, or has undergone removal or ablation of a colorectal neoplasia; and

Claims 5, 6, 12 and 13 should read as follows:

5. The method of claim 1, wherein the patient has undergone removal or ablation of a colorectal neoplasia.
6. The method of claim 1, wherein the aminoglycoside antibiotic is selected from the group consisting of: Amikacin, Gentamicin, Kanamycin, Neomycin, Netilmicin, Paromomycin, Streptomycin, and Tobramycin.
12. The method of claim 1, wherein the delivering step is effected by delivering a constant dosage of the aminoglycoside.
13. The method of claim 1, wherein the delivering step is effected by delivering a varying dosage of the aminoglycoside.

MAILING ADDRESS OF SENDER:

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PATENT NO. 6,846,807

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This collection of information is required by 37 CFR 1.322, 1.323, and 1.324. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 1.0 hour to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Attention Certificate of Corrections Branch, Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

If you need assistance in completing the form, call 1-800-PTO-9199 and select option 2.

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Applicant's remarks filed on 06/24/04 in response to the Office Action dated 02/25/04
are acknowledged.

The rejection of claims 1-5 and 7-17, under 35 U.S.C., 112, second paragraph, has
been overcome through the examiner's amendments.

A Declaration under 37 C.F.R. 1.132, executed by Jonathan Michael Greff, the inventor
of the present application has been entered.

The examiner withdraws the 35 U.S.C. 103(a) rejections, as being unpatentable over
Flotte et al. in view of Hu et al. in response to applicant's remarks that "The Hu patent
does not suggest to those skilled in the art that such enterically delivered
aminoglycosides could provide effective therapy for colorectal neoplasia, and Flotte's
method for using high-pressure impulse transients to promote drug delivery does not
somehow complement Hu to provide such suggestion".

1. An examiner's amendment to the record appears below. Should the changes
and/or additions be unacceptable to applicant, an amendment may be filed as provided
by 37 CFR 1.312. To ensure consideration of such an amendment, it MUST be
submitted no later than the payment of the issue fee.

Authorization for this examiner's amendment was given by Richard Osman on
10/13/04.

(1) Claims 6 and 18-20 have been cancelled without prejudice.

(2) Claims 16 and 17 depend from claim 15.

(3) Claims 1,5,7,13 and 14 have been amended as following:

1. (Currently Amended) A method of reducing development of colorectal neoplasia in a patient
subject or predisposed to colorectal neoplasia, the method comprising the steps of:

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determining a patient is subject or predisposed to colorectal neoplasia, or has undergone removal or ablation of a colorectal neoplasia; and

enterically delivering into the gut of the person an effective amount of an aminoglycoside antibiotic having poor gut absorption, whereby the development of the colorectal neoplasia is reduced as compared with otherwise similar non-treated patients.

5. (Currently Amended) The method of claim 1, wherein the patient has undergone removal or ablation of a colorectal neoplasia and is determined to be predisposed to colorectal neoplasia recurrence.

7. (Currently Amended) The method of claim 1, wherein the aminoglycoside antibiotic is selected from the group consisting of: Amikacin (Amikin[®]), Gentamicin (Garamycin[®]), Kanamycin (Kantrex[®]), Neomycin (Fucidin[®]), Neilmicin (Netromycin[®]), Paromomycin (Humatin[®]), Streptomycin, and Tobramycin (TOBI Salution[®], TobraDex[®], Nebein[®]).

13. (Currently Amended) The method of claim 1, wherein the delivering step is effected by delivering a constant over-time dosage of the aminoglycoside.

14. (Currently Amended) The method of claim 1, wherein the delivering step is effected by delivering a varying over-time dosage of the aminoglycoside.

Claims 1-5 and 7-17 are currently pending in this application.

Claims 1-5 and 7-17 are allowed.

2. The following is an examiner's statement of reasons for allowance: The instant claims are directed to a method of reducing development of colorectal neoplasia in a patient subject or predisposed to colorectal neoplasia comprising the steps of : determining a patient is subject to colorectal neoplasia, or has undergone removal or ablation of a colorectal neoplasia; and enterically delivering into the gut of the person an effective amount of an aminoglycoside antibiotic having poor gut absorption, whereby the development of the colorectal neoplasia is reduced as compared with otherwise

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